In the Claims

- 1. (previously presented) A method for inducing an immune response to a feline immunodeficiency virus (FIV) in a human or a non-feline animal that is susceptible to infection by FIV, said method comprising administering an effective amount of an FIV immunogen to said human or non-feline animal to induce said immune response.
- 2. (original) The method according to claim 1, wherein said FIV immunogen induces a humoral immune response.
- 3. (original) The method according to claim 1, wherein said FIV immunogen induces a cellular immune response.
- 4. (previously presented) The method according to claim 1, wherein said FIV immunogen induces an immune response against more than one subtype of FIV.
- 5. (previously presented) The method according to claim 1, wherein said FIV immunogen is selected from the group consisting of synthetic FIV peptide, natural or recombinant FIV protein or a fragment thereof, polynucleotide comprising a sequence that encodes an FIV protein or fragment thereof, polynucleotide comprising a sequence that encodes an FIV protein or a fragment thereof and an HIV protein or a fragment thereof, inactivated or attenuated whole FIV viral isolate, FIV viral fragment, inactivated cells infected with FIV, and a composition comprising FIV and HIV proteins or fragments thereof.
- 6. (original) The method according to claim 5, wherein said FIV immunogen comprises an epitope of an FIV and HIV protein that is evolutionarily conserved between the viruses.

- 7. (original) The method according to claim 6, wherein said protein is selected from the group consisting of core gag protein and envelope protein.
- 8. (original) A method for inducing an immune response to a human immunodeficiency virus (HIV) in a human, said method comprising administering an effective amount of an FIV immunogen to said human to induce said immune response.
- 9. (original) The method according to claim 8, wherein said FIV immunogen induces a humoral immune response.
- 10. (original) The method according to claim 8, wherein said FIV immunogen induces a cellular immune response.
- 11. (previously presented) The method according to claim 8, wherein said FIV immunogen induces an immune response against more than one subtype of FIV.
- 12. (previously presented) The method according to claim 8, wherein said FIV immunogen is selected from the group consisting of synthetic FIV peptide, natural or recombinant FIV protein or a fragment thereof, polynucleotide comprising a sequence that encodes an FIV protein or a fragment thereof, polynucleotide comprising a sequence that encodes an FIV protein or a fragment thereof and an HIV protein or a fragment thereof, inactivated or attenuated whole FIV viral isolate, FIV viral fragment, inactivated cells infected with FIV, and a composition comprising FIV and HIV proteins or fragments thereof.
- 13. (original) The method according to claim 12, wherein said FIV immunogen comprises an epitope of an FIV and HIV protein that is evolutionarily conserved between the viruses.
- 14. (original) The method according to claim 13, wherein said protein is selected from the group consisting of core gag protein and envelope protein.

15-46. (canceled)

- 47. (new) The method according to claim 1, wherein said FIV immunogen induces a protective immune response.
- 48. (new) The method according to claim 1, whereby an immune response is induced against FIV.
- 49. (new) The method according to claim 8, wherein said FIV immunogen induces a protective immune response.
- 50. (new) The method according to claim 8, whereby an immune response is induced against HIV.